AMENDMENTS TO THE SPECIFICATION

Please amend the specification rewriting paragraph 0010 of the published application as follows:

The present invention provides that the carrier of the oral dosage formulation be substantially solid polyethylene glycol. In one aspect of the present invention, the polyethylene glycol used may have an average molecular weight of from about 100 to about 20,000 or a mixture thereof. In another aspect, the polyethylene glycol carrier may have an average molecular weight of from about 1,000 to about 10,000 or a mixture thereof. In another embodiment of the present invention, the polyethylene glycol carrier may have an average molecular weight of from about 400 to about 15,000. Furthermore, in some aspects, the oral dosage formulation may include an amount of substantially solid polyethylene glycol of from about 30% w/w to about 80% w/w of the oral dosage formulation. In another aspect, the amount may be from about 50% w/w to about 80% w/w of the oral dosage formulation. In a further aspect, the amount may be from about 60% w/w to about 80% w/w. In yet another aspect, the amount may be 70% w/w of the oral dosage formulation.